## **AMENDMENTS**

## IN THE CLAIMS

Please amend claims 1, 13-16, 27, 28, 31-33, 36, 48, and 49 as shown below. Please cancel claims 27-33 and 36 herein without prejudice to their renewal.

- 1. (Presently Amended) A method for treating subject having a hemoglobinopathy in a subject, the method comprising administering to the subject in need thereof a compound, wherein the compound that inhibits hypoxia-inducible factor (HIF) prolyl hydroxylase, and wherein the compound increases expression of the gene encoding γ-globin in a bone marrow-derived cell, a hematopoietic stem cell, or a blast-forming unit erythroid cell or population of bone marrow-derived cells, thereby treating the hemoglobinopathy in the subject.
- 2-11. (Previously Canceled)
- 12. (Previously Amended) The method of claim 1, wherein the hemoglobinopathy comprises an alteration in the level, structural integrity, or activity of adult β-globin.
- 13. (Presently Amended) The method of claim 1, wherein the hemoglobinopathy is selected from the group consisting of β-thalassemias and sickle cell syndromes.
- 14. (Presently Amended) The method of claim 13, wherein the  $\beta$ -thalassemia is selected from the group consisting of  $\beta^0$ -thalassemia and  $\beta^+$ -thalassemia.
- 15. (Presently Amended) The method of claim 13, wherein the sickle cell syndrome is selected from the group consisting of sickle trait, sickle β-thalassemia, and sickle cell anemia.
- 16. (Presently Amended) The method of claim 1, wherein <u>administering the compound increases</u> the proportion of fetal hemoglobin relative to non-fetal hemoglobin produced by the bone marrow-

derived cell, the hematopoietic stem cell, or the blast-forming unit erythroid cell or population of cells in the subject is increased.

- 17-26. (Previously Canceled)
- 27-33. (Canceled Herein)
- 34-35. (Previously Canceled)
- 36. (Canceled Herein)
- 37-47. (Previously Canceled)
- 48. (Presently Amended) The method of claim 1, wherein the <u>compound that inhibits</u> HIF prolyl hydroxylase <del>inhibitor</del> is selected from the group consisting of an iron chelator, a <u>structural</u> <u>mimetic of 2-oxoglutarate-mimetic</u>, and a proline analog.
- 49. (Presently Amended) The method of claim 48, wherein the <u>structural mimetic of 2-oxoglutarate</u> mimetic inhibits HIF prolyl hydroxylase competitively with respect to 2-oxoglutarate and noncompetitively with respect to iron.